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# Quarterly Technical Progress Report December 2014

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December 16, 2014

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This work performed under the auspices of the U.S. Department of Energy by Lawrence Livermore National Laboratory under Contract DE-AC52-07NA27344.

# Quarterly Technical Progress Report

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Award Number:	10567486
Log Number:	LC130820
Project Title:	Mechanism for Clastogenic Activity of Naphthalene
Principal Investigator Name:	Bruce Buchholz
Principal Investigator Organization and Address:	Lawrence Livermore National Laboratory P.O. Box 808, L-397 Livermore, CA 94551
Principal Investigator Phone and Email:	925-422-1739 buchholz2@llnl.gov
Report Date:	December 12, 2014
Report Period:	September 1, 2014-November 30, 2014

**LLNL-TR-665462**

This work performed under the auspices of the U.S. Department of Energy by Lawrence Livermore National Laboratory under Contract DE-AC52-07NA27344.

## 1. Accomplishments:

The project has two main goals: 1) Identify the types of adducts naphthalene (NA) forms with DNA and 2) determine whether adduct formation correlates with site selective tumor formation in defined subcompartments of the respiratory tract (respiratory and olfactory nasal epithelium and airways of mice, rats and rhesus monkeys). Five tasks are associated with the completion of the goals.

Task 1: Contracting and Animal Use Approvals. IACUC and ACURO approvals are complete, The subcontract with UC Davis (UCD) was sent to them in mid-November but had not been returned with signatures by the end of the quarter as planned.

Task 2: Perform In Vitro Study for Goal 1. Scheduled to start in November but awaiting completion of subcontract with UCD. Anticipate commencement in December 2014.

Task 3: Perform In Vitro Study for Goal 2. Scheduled to start in February 2015.

Task 4: Sample Preparation and Analysis. Scheduled to start in November. Waiting for Samples.

Task 5: Data Interpretation and Reporting. Waiting for Samples. Anticipate sufficient data to submit abstract to SOT by mid January 2015 deadline.

### What was accomplished under these goals?

The major activities of the quarter were getting protocol approvals in place with UCD IACUC and ACURO. The subcontract is awaiting signature at UCD. Preparation for the studies is complete.

### Describe the Regulatory Protocol and Activity Status (if applicable).

#### (a) Human Use Regulatory Protocols

**TOTAL PROTOCOLS:** No human subjects research will be performed to complete the Statement of Work.

#### (b) Use of Human Cadavers for Research Development Test & Evaluation (RDT&E), Education or Training

No human cadaver research will be performed to complete the Statement of Work.

#### (c) Animal Use Regulatory Protocols

**TOTAL PROTOCOL(S):** 1 animal use protocol is required to complete the Statement of Work.

**PROTOCOL (1 of 1 total):**

Protocol [ACURO Assigned Number]: LC130820

Title: Quantitation of DNA Adducts from Naphthalene

Target required for statistical significance: 36 mice, 18 rats

Target approved for statistical significance: 36 mice, 18 rats

**SUBMITTED TO AND APPROVED BY:**

*Submitted to USAMRMC 14-AUG-2014*

*Approved by USAMRMC (ACURO) 15-OCT-2014 by Bryan K. Ketzenberger, DVM, DACLAM  
IACUC 18172 (UC Davis) submitted by Protocol PI Alan Buckpitt was approved 15-MAY-2014*

**STATUS:**

*Awaiting UCD signature on subcontract.*

*No technical or logistical issues.*

**What do you plan to do during the next reporting period to accomplish the goals and objectives?**

We anticipate the subcontract will be signed in December 2014.

We plan to complete most of the animal exposures required Goals 1 and 2. Samples will be analyzed by AMS.

Data Analysis will begin and an abstract will be submitted to SOT.

**2. Products:**

Nothing to report.

**3. Participants & Other Collaborating Organizations**

No Personnel have worked 1 person month on the project.

Name: Bruce Buchholz

Project Role: PI

Nearest person month worked: 0

Contribution to Project: Completed paperwork for ACURO approval and subcontract with UCD

#### **4. Changes/Problems::**

##### **a. Actual Problems or delays and actions to resolve them**

The delay in UCD signing the subcontract is minor at this point. The PIs at UCD are contacting their administration to encourage completion of the contract so work can begin.

##### **b. Anticipated Problems/Issues**

None anticipated.

#### **5. Special Reporting Requirements:**

**Quad Charts:** Quad chart attached.

# Mechanism for Clastogenic Activity of Naphthalene

LC130820

MIPR:10567486

PI: Bruce Buchholz

Org: Lawrence Livermore National Laboratory

Award Amount: \$165K

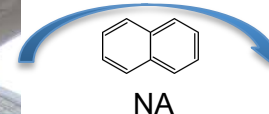


## Study/Product Aim(s)

- Identify the types of adducts naphthalene (NA) forms with DNA
- Determine whether adduct formation correlates with site selective tumor formation in defined subcompartments of the respiratory tract

## Approach

The location of lung and nasal tumors in rodents exposed to NA are highly specific to defined regions within the respiratory tract. We will utilize microdissection methods to isolate live tissue from these target areas. This approach combined with accelerator mass spectrometry (AMS) will determine whether DNA adducts are formed in the target tissue following incubation with  $^{14}\text{C}$ -NA.



Determine if naphthalene (NA) in jet fuel and cigarette smoke forms DNA adducts that can lead to cancer in respiratory tissues

Accomplishment: Regulatory approvals are in place and animal work will begin as soon as the subcontract is executed.

## Timeline and Cost

Activities	CY	14	15
Animal Protocol & Contract Complete		<div><div></div></div>	
In Vitro Studies for Aim 1 Complete		<div><div></div></div>	
In Vitro Studies for Aim 2 Complete		<div><div></div></div>	
Sample Analyses Complete		<div><div></div></div>	
Data Analyses and Reporting		<div><div></div></div>	
Estimated Budget (\$K)		\$15	\$150

## Goals/Milestones

**Q1 Goals** – Approval of Animal Protocol and Subcontract Executed

- ☒ Animal protocols approved
- ☐ Subcontract with UCD executed

**Q2 Goal** – Begin Experimental Work

- ☐ Begin in vitro studies and sample analyses for aims 1 & 2

**Q3 Goal** – Complete Experimental Work

- ☐ Finish all in vitro studies and sample analyses

**Q4 Goals** – Complete Data Analyses and Reporting

- ☐ Complete and submit peer-reviewed publication
- ☐ Complete and submit final report

## Comments/Challenges/Issues/Concerns

- Ready to begin in vitro studies as soon as subcontract is executed.

## Budget Expenditure to Date

Projected Expenditure: \$10K

Actual Expenditure: \$9K

Updated: (30-November-2014)